KANSAS DEPARTMENT OF CORRECTIONS

DOC Serving Konsas	Internal Management Policy and Procedure	SECTION NUMBER 06-101 SUBJECT: EVALUATION AND RES Evaluation Activities	PAGE NUMBER 1 of 6 EARCH: Research and
The wall		Original Date Issued:	08-15-82
		Replaces Amendment I	
Reissued By: Policy & Procedure Coordinator		The substantive content of this IMPP has been reissued as per the appropriate provisions of IMPP 01-101. The only modifications within the reissue of this document concern technical revisions of a non substantive nature. Date Reissued: 05-20-11	

POLICY

Research and evaluation activities by individuals and/or organizations outside the Department are permitted and encouraged, if they are relevant to the Department and its programs. (ACO 2-1F-10, 2-1F-11, ACI 3-4105, 3-4106, APPFS 2-3096) System Management Team members are encouraged and expected to undertake and/or support research and evaluation activities to assess the efficiency and effectiveness of operations, programs and/or services under their management. (ACO 2-1F-10, 2-1F-11, ACI 3-4105, 3-4106, APPFS 2-3096) System Management Team members may independently conduct and/or authorize the conduct of research and evaluation activities, (ACO 2-1F-12, ACI 3-4109) provided the results of such activities will not be submitted for publication or other professional/academic distribution and the authorizing System Management Team member is not cited as one of the researchers. Proposals for research projects which may possibly be submitted for publication or other professional or academic distribution shall be submitted for review by designated Central Office staff in advance to ensure that appropriate methodologies and procedures are used; such proposals shall be subject to final approval by the Secretary. Advance review by designated Central Office staff and approval by the Secretary shall be obtained prior to contracting for any research activity, regardless of how such research will be utilized or distributed. Appropriate safeguards and limitations shall be utilized to protect the welfare and privacy of individual staff and offenders involved as subjects. (ACO 2-1F-12, 2-1F-15, APPFS 2-3101)

Research and evaluation activities shall be conducted and the results reported, disseminated and utilized in conformity with Department policies and procedures, professional and scientific ethics, and, with State and Federal guidelines for the use and dissemination of research findings. (ACO 2-1F-09, ACI 3-4108 NCCHC P-72) All completed research or evaluation reports shall be submitted for review by the Management Team prior to being submitted for publication or otherwise released for distribution.

Offenders shall not be used for medical, pharmaceutical or cosmetic research, or, experiments. This prohibition shall not preclude any individual offender from receiving treatment via a specific medical procedure that is not generally available when such treatment procedure is approved by the appropriate governmental agencies, Departmental Health Authority, and the offender's consent is documented. (ACO 2-1F-14, ACI 3-4373) Participation in biomedical, non-medical, non-pharmaceutical and non-cosmetic research, including sociological and psychological research involving human subjects shall require the written informed, voluntary consent of the staff and offenders involve d. (ACO 2-1F-13, ACI 3-4110, NCCHC P-72)

DEFINITIONS

<u>Data</u>: Questionnaires, inventories, and aggregated information on hard copy or computer diskettes.

<u>Departmental Health Authority</u>: The medical director of the agency or organization responsible for the provision of health care for the Kansas Department of Corrections.

<u>Facility health authority</u>: The physician or health administrator responsible for the provision of health care services at a facility. The facility health authority works under direction of the Department's health authority.

<u>In-house research/evaluation</u>: Activities or projects initiated by a System Management Team member to assess the efficiency or effectiveness of any aspect of their operation.

<u>Management Team (MT)</u>: A panel of Central Office management staff designated by the Secretary. Currently this panel is comprised of the Secretary; deputy secretaries; chief legal counsel; staff assistant to the Secretary; public information officer; human resources manager; information resource manager; and the fiscal officer.

<u>Principal Administrator</u>: The person directly responsible for the overall administration of a KDOC facility, parole region, or Central Office work unit.

<u>System Management Team (SMT)</u>: A management panel designated by the Secretary which is comprised of the Management Team, wardens, parole directors, and Kansas Correctional Industries directors.

<u>User</u>: The person, agency or group authorized to engage in research activity within a facility, office or unit of the Department of Corrections.

PROCEDURES

I. Encouragement and Facilitation of Research Activities

- A. To encourage and facilitate research activities, the Department shall seek funding for research projects relevant to stated goals and objectives.
- B. In all cases where research is being proposed by persons not employed by the Department, or those persons employed by the Department in a position where research activity is not a part of the person's job description, an Access Request and Non-Disclosure Agreement (hereinafter referred to as "user's agreement") shall be completed and submitted with the research proposal. See Attachment A. form 06-101-001.

II. Provision for Treatment and Prohibition of Certain Research Activities

- A. When a specific medical procedure not generally available is recommended in the treatment of an individual offender, such treatment shall not begin until after a full explanation of the positive and negative features of the treatment has been given to the offender.
 - The explanation to the offender shall be documented, in accordance with procedures established in IMPP 10-127, utilizing the Consent to Treat form for surgical and nonroutine medical treatment.
- B. Except as provided above, all research proposals involving offender participation in medical, pharmaceutical or cosmetic testing for experimental research purposes shall be rejected.

III. Voluntary Informed Consent Agreement

A. Research involving human subjects, whether employees or offenders, shall require the documented, voluntary informed consent of each participant. Such consent shall utilize appropriate forms, as provided in IMPP 10-127, and shall be obtained in advance of the subjects participation.

- B. At a minimum, obtaining an informed consent shall include:
 - 1. A fair explanation of the procedures to be followed including an identification of that which is experimental;
 - 2. A description of the potential discomforts and/or risks;
 - 3. A description of the benefits to be expected;
 - 4. A description of appropriate alternative procedures;
 - 5. An offer to answer any inquiries concerning proposed procedures; and,
 - 6. Instruction that the subject is free to withdraw consent and to discontinue participation in the project at any time.
- C. Use of subjects who are legally unable to give the informed consent (e.g., under the age of 18 or mentally incompetent) shall be prohibited. (ACO 2-4E-01, ACI 3-4372, NCCHC P-70)

IV. Confidentiality of Research Data

- A. Provisions shall be made by the user for safeguarding the confidentiality and prohibiting the dissemination of research data that can be traced to or identified with individual subjects. (ACO 2-1F-15)
- B. Questionnaires, inventories and other data gathering instruments and/or procedures shall limit identifying and/or personal information recorded to only that essential to the project as specified within the context of the user's agreement.
 - 1. Research data containing information from which the identity of subjects can be traced shall not be disseminated to anyone except appropriate project or KDOC staff.
- C. All data collected by a KDOC employee as part of the duties or activities of the position shall be the property of the facility, parole district, or unit in which the data was collected and maintained in accordance with the records retention schedule established by the State Records Board.
- D. Data collected by an outside researcher or by a KDOC employee who, while off duty, conducts research that is not a part of his/her position responsibility shall be considered the property of that researcher or the agency the researcher represents. Such data shall be maintained in accordance with guidelines established by the agency represented by the researcher.

V. Review and Approval Process for Research and Evaluation Proposals

- A. System Management Team members conducting or authorizing in-house research or evaluation projects shall be responsible to ensure the research design and methodologies are in conformance with the provisions of this IMPP. (ACI 3-4109)
- B. Except for in-house research and evaluation projects approved by System Management Team members, all proposals for research studies shall be forwarded to the Research and Planning Unit for review prior to the initiation of any research activity. (ACO 2-1F-12)
 - 1. Research/evaluation proposals shall be submitted to the Research and Planning Unit when one or more of the following conditions exist:

- a. The research or assessment to be conducted is not under the direction of a System Management Team member.
- b. The research or findings will be submitted for publication in a newspaper, newsletter, magazine or professional journal or in a paper submitted in conjunction with a college/university program or presented at a professional conference or meeting.
- The System Management Team member of the organizational unit is or will be cited as one of the researchers.
- C. Research proposals shall address all of the design and methodology issues and include all information and material listed in the Research Proposal Format, Attachment B.
- D. Each member of the Management Team shall be given notice of any research proposals that are received by the Research and Planning_Unit for review and each such person shall be given the option of reviewing each proposal, consistent with his/her interests.
 - 1. The principal administrator at each facility/region/unit involved in the research shall be given a copy of the full proposal to review.
- E. The Research and Planning Unit shall determine:
 - 1. The potential benefits to science, society and the intended subjects;
 - 2. The potential risks and costs to the intended subjects; and,
 - 3. A recommended course of action, based upon a professional assessment of the anticipated benefits to inmates, the Department, and the Secretary of Corrections.
- F. Only those proposed projects wherein potential benefits clearly outweigh potential risks and costs shall be given any further consideration.
- G. Those projects not meeting minimum approval criteria as established under V. E. and not receiving further consideration under V. F. shall be recommended for denial.
- H. For those projects meriting further consideration, the Research and Planning Unit shall conduct a review to determine:
 - 1. The relative merit and appropriateness of the project, i.e., consistency with the KDOC Mission statement and relevance to the Department's programs, services and operations; (ACO 2-1F-10, ACI 3-4105, APPFS 2-3096)
 - 2. The qualifications of the researcher(s);
 - 3. The adequacy of:
 - a. The research design;
 - b. The voluntary informed consent agreement; and,
 - c. The procedures designed to maintain the confidentiality, security and privacy of research data. (ACO 2-1F-15)
 - 4. The disruptive effects, if any, upon the orderly management of the project site; and,

- 5. Such input from principal administrators as can be obtained and considered during the course of the proposal review.
- Within ten (10) days of receipt of the research proposal, the Research and Planning Unit shall document its review and forward the review to the Secretary of Corrections for approval or disapproval.
- J. Within ten (10) days of receipt, the Secretary of Corrections shall render a written decision concerning the proposal which shall:
 - 1. Permit the research to proceed;
 - 2. Make alterations, or, request that alterations be made in the proposal and the proposal be resubmitted; or,
 - 3. Deny permission for the research. (APPFS 2-3098)
- K. The researcher shall be informed of the Secretary's decision in writing by the Research and Planning Unit Manager or designee. A copy of the notification letter shall be forwarded to the principal administrator of the facility/region/unit involved in the research by the Research and Planning Unit.
- L. Researchers seeking a reconsideration of the Secretary of Corrections' decision may contact the Secretary of Corrections or designee for further discussion and review of the project.

VI. Response to Possible User's Agreement Violations

- A. Upon receipt of sufficient information indicating a violation of the user's agreement, the principal administrator shall:
 - 1. Suspend the activities of the research project; and,
 - 2. Notify the Chief Legal Counsel, who shall initiate and direct further action taken on the matter.
- B. If the Chief Legal Counsel determines that violation of the user's agreement has occurred, the following action shall be taken:
 - 1. The project shall be terminated;
 - 2. Data collected shall be confiscated and submitted to the records section for storage; and,
 - 3. Appropriate sanctions as listed in the user's agreement shall be applied.
- C. If the Chief Legal Counsel determines that no violation of the user's agreement has occurred:
 - 1. The appropriate principal administrator shall be so notified; and,
 - 2. Upon such notification, the principal administrator shall rescind the suspension of research activities effected under VI. A. 1., above.

VII. Reporting Process for Approved Research Proposals and Dissemination of Findings

- A. A final written report on all research or evaluation activities must be submitted to the Research and Planning Unit no later than sixty (60) days after the completion of the activity and prior to being submitted for publication or other release for distribution.
 - 1. The final report shall be reviewed by the Research and Planning Unit, forwarded to the Secretary of Corrections and the appropriate principal administrator(s) for review. (ACO 2-1F-04, APPFS 2-3102)

VIII. Publication of Completed Research

- A. Completed research or evaluation projects may be submitted by the author(s) to professional journals for publication, with the approval of the Secretary of Corrections or designee.
- B. Research or evaluation projects which, upon completion, are not submitted for publication in professional journals may be compiled and published by the Department on a regular basis with the approval of the Secretary of Corrections or designee.

IX. Disclaimer Requirement

A. All manuscripts prepared in an unofficial capacity and submitted for publication by departmental employees, including employees of entities with which the Department has contractual arrangements, shall contain a disclaimer which states that any conclusions, interpretations or recommendations expressed in the manuscript are those of the author and do not necessarily reflect the position or policy of the Kansas Department of Corrections.

NOTE: The policy and procedures set forth herein are intended to establish directives and guidelines for staff and offenders and those entities that are contractually bound to adhere to them. They are not intended to establish State created liberty interests for employees or offenders, or an independent duty owed by the Department of Corrections to employees, offenders, or third parties. Similarly, those references to the standards of various accrediting entities as may be contained within this document are included solely to manifest the commonality of purpose and direction as shared by the content of the document and the content of the referenced standards. Any such references within this document neither imply accredited status by a Departmental facility or organizational unit, nor indicate compliance with the standards so cited. The policy and procedures contained within this document are intended to be compliant with all applicable statutes and/or regulatory requirements of the Federal Government and the state of Kansas. This policy and procedure is not intended to establish or create new constitutional rights or to enlarge or expand upon existing constitutional rights or duties.

REPORTS REQUIRED

Name/Type of Report	By Whom/To Whom	<u>Due</u>
Final Project Report	Researcher to Research and Planning Unit	Within 60 days after completion of project

<u>REFERENCES</u>

IMPP 10-127
ACO 2-1E-01, 2-1F-04, 2-1F-09, 2-1F-10, 2-1F-11, 2-1F-12, 2-1F-13, 2-1F-14, 2-1F-15
ACI 3-4105, 3-4106, 3-4108, 3-4109, 3-4110, 3-4372, 3-4373
APPFS 2-3096, 2-3098, 2-3101, 2-3102
NCCHC P-70, P-72

ATTACHMENTS

Attachment A - Access Request and Non-disclosure Agreement, 1 page Attachment B - Research Proposal Format, 2 pages

ACCESS REQUEST AND NON-DISCLOSURE AGREEMENT FOR INFORMATION PERTAINING TO INMATES IN THE KANSAS CORRECTIONAL SYSTEM

	agreement sets forth conditions under which access to selected inmate information will be provided by the as Department of Corrections to,				
	, hereinafter called				
Requ	estor.				
1.	Information Requested:				
2.	Requestor requests this information () on a continuing basis () on a one-time basis				
3.	The purpose for which information requested is: (check one)				
()	To implement a statute or executive order that expressly refers to criminal conduct and contains requirements and/or exclusions expressly based upon such conduct. Give citation:				
()	To carry out a contract or agreement to provide services required for the administration of justice. Attac agreement.				
()	To conduct research, evaluative, or statistical activities.				
()	To implement a state or federal statute or executive order to conduct investigations determining employment suitability or eligibility for security clearances allowing access to classified information pursuant to a state or federal statute or executive order. Give citation:				
()	To exercise authority granted by court order or rule. Attach order or rule.				
()	Other purpose, as described below or in attachment.				
4.	Requestor agrees to limit the use of any received information to the purpose(s) for which it was provided and to destroy the information when it is no longer needed for the purpose(s) for which it was provided.				
5.	Requestor agrees that the only person(s) allowed access to any received information are those name here; and to not disseminate the information to any other agency or person:				
Requ	estor:				
Name	e				
Agen Title_	cy				
Signa	ature Date				
Kans	as Department of Corrections:				
Name					
Title					
Signa	ature Date				

Form # 06-101-001

RESEARCH PROPOSAL FORMAT

- I. Title of Study
 - A. Name of Author(s)
 - Institutional Affiliation
 - Qualifications
- II. Timetable for the study, including estimated dates of implementation and completion.
- III. Personnel needs, indicating the time to be spent by each, and the availability of each.
- IV. Materials needed for the project, and whether such material is available or will need to be secured.
- V. Project Design
 - A. Introduction
 - 1. Present a clear, concise statement of the research problem and why it is worthy of study.
 - 2. Review the literature by briefly summarizing the findings from other research which is relevant to the research problem.
 - Describe the purpose of the study.
 - State the hypotheses of the study.
 - 5. Identify the factors whose effects are to be studied (independent variables) and the factors on which measures will be taken (dependent variables).
 - a. Explain any proposed manipulations of independent variables (identification of any experimental treatment to be imposed).
 - b. State precisely how the dependent variable will be measured.
 - c. Explain any procedures that will be implemented to control for other variables that could intervene.

B. Method

- 1. Subjects: Identify the research subjects or study groups and describe their demographic characteristics.
 - a. Submit voluntary informed consent agreement.
 - b. Describe and attach any experimental apparatus, survey instruments, or testing instruments to be employed in the study.
 - c. Describe concisely and exactly what will be required of the participant(s); how experimental sessions with the subject(s) will be conducted; and, by whom or how questionnaires or tests will be administered.

d. Proposed Data Analysis

- (1) Describe the form in which the data will be collected and exactly how data will be analyzed. Include a description of statistical testing to be performed.
- (2) Discuss what results would support the hypotheses, and what results would refute the hypotheses.